Dear Ms. Garcia,

The Division of Health Improvement/Quality Management Bureau has completed a compliance review survey of the services identified above. The purpose of the survey was to determine compliance with federal and state standards; to assure the health, safety, and welfare of individuals receiving services through the Developmental Disabilities Waiver; and to identify opportunities for improvement.

**Quality Management Determination of Compliance:**
The Division of Health Improvement is issuing your agency a determination of “Non-Compliance with Conditions of Participation.”

**Plan of Correction:**
The attached Report of Findings identifies deficiencies found during your agency’s survey. You are required to complete and implement a Plan of Correction (POC). Please submit your agency’s Plan of Correction (POC) in the space on the two right columns of the Report of Findings. See attachment A for additional guidance in completing the POC. The response is due to the parties below within 10 working days of the receipt of this letter:

1. Quality Management Bureau, Attention: Plan of Correction Coordinator
   5301 Central Ave. NE Suite 400 Albuquerque, NM 87108

---

“Assuring safety and quality of care in New Mexico’s health facilities and community-based programs.”

David Rodriguez, Division Director • Division of Health Improvement
Quality Management Bureau • 5301 Central Ave. NE Suite 400 • Albuquerque, New Mexico 87108
(505) 222-8623 • FAX: (505) 222-8661 • http://dhi.health.state.nm.us
Upon notification from QMB that your Plan of Correction has been approved, you must implement all remedies and corrective actions within 45 working days. If your plan of correction is denied, you must resubmit a revised plan ASAP for approval. All remedies must still be completed within 45 working days of the original submission.

Failure to submit, complete or implement your POC within the required time frames will result in the imposition of a $200 per day Civil Monetary Penalty until it is received, completed and/or implemented.

**Request for Informal Reconsideration of Findings (IRF):**
If you disagree with a determination of noncompliance (finding) you have 10 working days upon receipt of this notice to request an IRF. Submit your request for an IRF in writing to:

QMB Deputy Bureau Chief  
5301 Central Ave NE Suite #400  
Albuquerque, NM 87108  
Attention: IRF request

A request for an IRF will not delay the implementation of your Plan of Correction which must be completed within 45 working days. Providers may not appeal the nature or interpretation of the standard or regulation, the team composition, sampling methodology or the Scope and Severity of the finding.

If the IRF approves the change or removal of a finding, you will be advised of any changes.

Please call the Team Leader at 505-670-6290, if you have questions about the survey or the report. Thank you for your cooperation and for the work you perform.

Sincerely,

*Nadine Romero, LBSW*

Nadine Romero, LBSW  
Team Lead/Healthcare Surveyor  
Division of Health Improvement  
Quality Management Bureau
**Survey Process Employed:**

Entrance Conference Date: July 12, 2010

Present:

- **Salas Care, Inc.**
  - Jeanette Salas, Vice President/Service Coordinator

- **DOH/DHI/QMB**
  - Nadine Romero, LBSW, Team Lead/Healthcare Surveyor
  - Florie Alire, RN, Healthcare Surveyor

- **DDSD – Metro Regional Office**
  - Rose Mary Williams, Community Living Coordinator

Exit Conference Date: July 16, 2010

Present:

- **Salas Care, Inc.**
  - Margaret Garcia, President
  - Jeanette Salas, Vice President/Service Coordinator

- **DOH/DHI/QMB**
  - Nadine Romero, LBSW, Team Lead/Healthcare Surveyor
  - Florie Alire, RN, Healthcare Surveyor

Homes Visited Number: 1

Administrative Locations Visited Number: 1

Total Sample Size Number: 3

- 0 - Jackson Class Members
- 3 - Non-Jackson Class Members
- 3 - Supported Living
- 1 - Adult Habilitation

Persons Served Interviewed Number: 3

Records Reviewed (Persons Served) Number: 3

Administrative Files Reviewed

- Billing Records
- Medical Records
- Incident Management Records
- Personnel Files
- Training Records
- Agency Policy and Procedure
- Caregiver Criminal History Screening Records
- Employee Abuse Registry
- Human Rights Notes and/or Meeting Minutes
- Nursing personnel files
- Evacuation Drills
- Quality Improvement/Quality Assurance Plan

**CC:** Distribution List: DOH - Division of Health Improvement

DOH - Developmental Disabilities Supports Division

DOH - Office of Internal Audit

HSD - Medical Assistance Division
Attachment A

Provider Instructions for Completing the QMB Plan of Correction (POC) Process

• After a QMB Quality Review, your Survey Report will be sent to you via certified mail. You may request that it also be sent to you electronically by calling George Perrault, Plan of Correction Coordinator at 505-222-8647.
• Within 10 business days of the date you received your survey report, you must develop and send your Plan of Correction response to the QMB office. (Providers who do not pick up their mail will be referred to the Internal Review Committee [IRC]).
• For each Deficiency in your Survey Report, include specific information about HOW you will correct each Deficiency, WHO will fix each Deficiency (“Responsible Party”), and by WHEN (“Date Due”).
• Your POC must not only address HOW, WHO and WHEN each Deficiency will be corrected, but must also address overall systemic issues to prevent the Deficiency from reoccurring, i.e., Quality Assurance (QA). Your description of your QA must include specifics about your self-auditing processes, such as HOW OFTEN you will self-audit, WHO will do it, and WHAT FORMS will be used.
• Corrective actions should be incorporated into your agency’s Quality Assurance/Quality Improvement policies and procedures.
• You may send your POC response electronically to George.Perrault@state.nm.us, by fax (505-222-8661), or by postal mail.
• Do not send supporting documentation to QMB until after your POC has been approved by QMB.
• QMB will notify you if your POC has been “Approved” or “Denied”.
• Whether your POC is “Approved” or “Denied”, you have a maximum of 45 business days to correct all survey Deficiencies from the date of receipt of your Survey Report. If your POC is “Denied” it must be revised and resubmitted ASAP, as the 45 working day limit is in effect. Providers whose revised POC is denied will be referred to the IRC.
• The POC must be completed on the official QMB Survey Report and Plan of Correction Form, unless approved in advance by the POC Coordinator.
• If you have questions about the POC process, call the QMB POC Coordinator, George Perrault at 505-222-8647 for assistance.
• For Technical Assistance (TA) in developing or implementing your POC, contact your local DDSD Regional Office.
• Once your POC has been approved by QMB, the POC may not be altered or the dates changed.
• Requests for an extension or modification of your POC (post approval) must be made in writing and submitted to the POC Coordinator at QMB, and are approved on a case-by-case basis.
• When submitting supporting documentation, organize your documents by Tag #s, and annotate or label each document using Individual numbers.
• Do not submit original documents, hard copies or scanned and electronically submitted copies are fine. Originals must be maintained in the agency/client file(s) as per DDSD Standards.
• Failure to submit, complete or implement your POC within the required timeframes will result in a referral to the IRC and the possible imposition of a $200 per day Civil Monetary Penalty until it is received, completed and/or implemented.
QMB Scope and Severity Matrix of survey results

Each deficiency in your Report of Findings is scored on a Scope and Severity Scale. The culmination of each deficiency's Scope and Severity is used to determine degree of compliance to standards and regulations and level of QMB Certification.

<table>
<thead>
<tr>
<th>SEVERITY</th>
<th>SCOPE</th>
<th>Isolated 01% - 15%</th>
<th>Pattern 16% - 79%</th>
<th>Widespread 80% - 100%</th>
</tr>
</thead>
<tbody>
<tr>
<td>Low Impact</td>
<td>No Actual Harm Minimal potential for harm.</td>
<td>A.</td>
<td>B.</td>
<td>C.</td>
</tr>
<tr>
<td>Medium Impact</td>
<td>No Actual Harm Potential for more than minimal harm</td>
<td>D. (2 or less)</td>
<td>E.</td>
<td>F. (3 or more)</td>
</tr>
<tr>
<td>High Impact</td>
<td>Actual harm</td>
<td>G.</td>
<td>H.</td>
<td>I.</td>
</tr>
<tr>
<td></td>
<td>Immediate Jeopardy to individual health and or safety</td>
<td>J.</td>
<td>K.</td>
<td>L.</td>
</tr>
</tbody>
</table>

Scope and Severity Definitions:

**Key to Scope scale:**

Isolated:
A deficiency that is limited to 1% to 15% of the sample, usually impacting no more than one or two individuals in the sample.

Pattern:
A deficiency that impacts a number or group of individuals from 16% to 79% of the sample is defined as a pattern finding. Pattern findings suggest the need for system wide corrective actions.

Widespread:
A deficiency that impacts most or all (80% to 100%) of the individuals in the sample is defined as widespread or pervasive. Widespread findings suggest the need for system wide corrective actions as well as the need to implement a Continuous Quality Improvement process to improve or build infrastructure. Widespread findings must be referred to the Internal Review Committee for review and possible actions or sanctions.

**Key to Findings:**

"Substantial Compliance with Conditions of Participation"
The QMB determination of “Substantial Compliance with Conditions of Participation” indicates that a provider is in substantial compliance with all ‘Conditions of Participation’ and other standards and regulations. The agency has obtained a level of compliance such that there is a minimal potential for harm to individuals’ health and safety. To qualify for a determination of Substantial Compliance with Conditions of Participation, the provider must not have any findings that meet the thresholds for determining non-compliance with any Condition of Participation.

"Non-Compliance with Conditions of Participation"
The QMB determination of “Non-Compliance with Conditions of Participation” indicates that a provider is out of compliance with one (1) or more ‘Conditions of Participation.’ This non-compliance, if not corrected, is likely to result in a serious negative outcome or the potential for more than minimal harm to individuals’ health and safety.

Providers receiving a repeat determination of Non-Compliance may be referred by QMB to the Internal Review Committee (IRC) for consideration of remedies and possible actions.

"Sub-Standard Compliance with Conditions of Participation”:
The QMB determination of “Sub-Standard Compliance with Conditions of Participation” indicates a provider is significantly out of compliance with Conditions of Participation and/or has:
Multiple findings of widespread non-compliance with any standard or regulation with a significant potential for more than minimal harm. Any finding of actual harm or Immediate Jeopardy.

Providers receiving a repeat determination of ‘Substandard Compliance’ will be referred by QMB to the Internal Review Committee (IRC) for consideration of remedies and possible actions.
Informal Reconsideration of Finding (IRF) Process

Introduction:
Throughout the process, surveyors are openly communicating with providers. Open communication means that surveyors have clarified issues and/or requested missing information before completing the review. Regardless, there may still be instances where the provider disagrees with a specific finding.

To informally dispute a finding the provider must request in writing an Informal Reconsideration of the Finding (IRF) to the QMB Deputy Bureau Chief within 10 working days of receipt of the final report.

The written request for an IRF must be completed on the QMB Request for Informal Reconsideration of Finding Form (available on the QMB website: http://dhi.health.state.nm.us/qmb) and must specify in detail the request for reconsideration and why the finding is inaccurate. The IRF request must include all supporting documentation or evidence that was not previously reviewed during the survey process.

The following limitations apply to the IRF process:

- The request for an IRF and all supporting evidence must be received in 10 days.
- Findings based on evidence requested during the survey and not provided may not be subject to reconsideration.
- The supporting documentation must be new evidence not previously reviewed by the survey team.
- Providers must continue to complete their plan of correction during the IRF process.
- Providers may not request an IRF to challenge the Scope and Severity of a finding.
- Providers may not request an IRF to challenge the sampling methodology.
- Providers may not request an IRF based on disagreement with the nature of the standard or regulation.
- Providers may not request an IRF to challenge the team composition.
- Providers may not request an IRF to challenge the QMB Quality Approval Rating and the length of their DDSD provider contract.

A Provider forfeits the right to an IRF if the request is not made within 10 working days of receiving the report and does not include all supporting documentation or evidence to show compliance with the standards and regulations.

QMB has 30 working days to complete the review and notify the provider of the decision. The request will be reviewed by the IRF committee. The Provider will be notified in writing of the ruling, no face to face meeting will be conducted.

When a Provider requests that a finding be reconsidered, it does not stop or delay the Plan of Correction process. Providers must continue to complete the Plan of Correction, including the finding in dispute regardless of the IRF status. If a finding is successfully reconsidered, it will be noted and will be removed or modified from the report. It should be noted that in some cases a Plan of Correction may be completed prior to the IRF process being completed. The provider will be notified in writing on the decisions of the IRF committee.
<table>
<thead>
<tr>
<th>Statute</th>
<th>Deficiency</th>
<th>Agency Plan of Correction and Responsible Party</th>
<th>Date Due</th>
</tr>
</thead>
<tbody>
<tr>
<td>Tag # 1A20 DSP Training Documents</td>
<td>Scope and Severity Rating: E</td>
<td>Based on record review, the Agency failed to ensure that Orientation and Training requirements were met for 6 of 12 Direct Service Personnel.</td>
<td></td>
</tr>
<tr>
<td>Developmental Disabilities (DD) Waiver Service Standards effective 4/1/2007</td>
<td></td>
<td>Review of Direct Service Personnel training records found no evidence of the following required DOH/DDSD trainings and certification being completed:</td>
<td></td>
</tr>
<tr>
<td>CHAPTER 1 IV. GENERAL REQUIREMENTS FOR PROVIDER AGENCY SERVICE PERSONNEL:</td>
<td></td>
<td>• Person-Centered Planning (1-Day) (DSP #42)</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Rights &amp; Advocacy (DSP #45)</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Teaching &amp; Support Strategies (DSP #43, 44, 45, 48 &amp; 51)</td>
<td></td>
</tr>
</tbody>
</table>
Developmental Disabilities Supports Division (DDSD) Policy - Policy Title: Training Requirements for Direct Service Agency Staff Policy - Eff. March 1, 2007 - II. POLICY STATEMENTS:

A. Individuals shall receive services from competent and qualified staff.
B. Staff shall complete individual-specific (formerly known as “Addendum B”) training requirements in accordance with the specifications described in the individual service plan (ISP) of each individual served.
C. Staff shall complete training on DOH-approved incident reporting procedures in accordance with 7 NMAC 1.13.
D. Staff providing direct services shall complete training in universal precautions on an annual basis. The training materials shall meet Occupational Safety and Health Administration (OSHA) requirements.
E. Staff providing direct services shall maintain certification in first aid and CPR. The training materials shall meet OSHA requirements/guidelines.
F. Staff who may be exposed to hazardous chemicals shall complete relevant training in accordance with OSHA requirements.
G. Staff shall be certified in a DDSD-approved behavioral intervention system (e.g., Mandt, CPI) before using physical restraint techniques. Staff members providing direct services shall maintain certification in a DDSD-approved behavioral intervention system if an individual they support has a behavioral crisis plan that includes the use of physical restraint techniques.
H. Staff shall complete and maintain certification in a DDSD-approved medication course in accordance with the DDSD Medication Delivery Policy M-001.
I. Staff providing direct services shall complete safety training within the first thirty (30) days of employment and before working alone with an individual receiving services.
<table>
<thead>
<tr>
<th>Tag # 1A26 (CoP) COR / EAR</th>
<th>Scope and Severity Rating: D</th>
</tr>
</thead>
<tbody>
<tr>
<td>NMAC 7.1.12.8</td>
<td>Based on record review, the Agency failed to maintain documentation in the employee’s personnel records that evidenced inquiry to the Employee Abuse Registry prior to employment for 2 of 13 Agency Personnel.</td>
</tr>
<tr>
<td>REGISTRY ESTABLISHED; PROVIDER INQUIRY REQUIRED:</td>
<td>The following Agency Personnel records contained evidence that indicated the Employee Abuse Registry was completed after hire:</td>
</tr>
<tr>
<td>Upon the effective date of this rule, the department has established and maintains an accurate and complete electronic registry that contains the name, date of birth, address, social security number, and other appropriate identifying information of all persons who, while employed by a provider, have been determined by the department, as a result of an investigation of a complaint, to have engaged in a substantiated registry-referred incident of abuse, neglect or exploitation of a person receiving care or services from a provider.</td>
<td></td>
</tr>
<tr>
<td>Additions and updates to the registry shall be posted no later than two (2) business days following receipt. Only department staff designated by the custodian may access, maintain and update the data in the registry.</td>
<td></td>
</tr>
<tr>
<td>A. <strong>Provider requirement to inquire of registry.</strong> A provider, prior to employing or contracting with an employee, shall inquire of the registry whether the individual under consideration for employment or contracting is listed on the registry.</td>
<td></td>
</tr>
<tr>
<td>B. <strong>Prohibited employment.</strong> A provider may not employ or contract with an individual to be an employee if the individual is listed on the registry as having a substantiated registry-referred incident of abuse, neglect or exploitation of a person receiving care or services from a provider.</td>
<td></td>
</tr>
<tr>
<td>D. <strong>Documentation of inquiry to registry.</strong> The provider shall maintain documentation in the employee’s personnel or employment records that evidences the fact that the provider made an inquiry to the registry concerning that employee prior to employment. Such documentation must include evidence, based on the response to such inquiry received from the custodian by the provider, that the employee was not listed on the registry as having a substantiated registry-referred incident of abuse, neglect or exploitation.</td>
<td></td>
</tr>
</tbody>
</table>

Based on record review, the Agency failed to maintain documentation in the employee’s personnel records that evidenced inquiry to the Employee Abuse Registry prior to employment for 2 of 13 Agency Personnel.

The following Agency Personnel records contained evidence that indicated the Employee Abuse Registry was completed after hire:

- #44 – Date of Hire 7/05/09
- #48 – Date of Hire 6/3/09
E. **Documentation for other staff.** With respect to all employed or contracted individuals providing direct care who are licensed health care professionals or certified nurse aides, the provider shall maintain documentation reflecting the individual's current licensure as a health care professional or current certification as a nurse aide.

F. **Consequences of noncompliance.** The department or other governmental agency having regulatory enforcement authority over a provider may sanction a provider in accordance with applicable law if the provider fails to make an appropriate and timely inquiry of the registry, or fails to maintain evidence of such inquiry, in connection with the hiring or contracting of an employee; or for employing or contracting any person to work as an employee who is listed on the registry. Such sanctions may include a directed plan of correction, civil monetary penalty not to exceed five thousand dollars ($5000) per instance, or termination or non-renewal of any contract with the department or other governmental agency.


**Chapter 1.IV. General Provider Requirements.**

**D. Criminal History Screening:** All personnel shall be screened by the Provider Agency in regard to the employee's qualifications, references, and employment history, prior to employment. All Provider Agencies shall comply with the Criminal Records Screening for Caregivers 7.1.12 NMAC and Employee Abuse Registry 7.1.12 NMAC as required by the Department of Health, Division of Health Improvement.
### Tag # 6L14  Residential Case File

<table>
<thead>
<tr>
<th>Developmental Disabilities (DD) Waiver Service Standards effective 4/1/2007</th>
<th>Scope and Severity Rating: D</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>CHAPTER 6. VIII. COMMUNITY LIVING SERVICE PROVIDER AGENCY REQUIREMENTS</strong></td>
<td>Based on record review, the Agency failed to maintain a complete and confidential case file in the residence for 1 of 3 Individuals receiving Supported Living Services.</td>
</tr>
<tr>
<td><strong>A. Residence Case File:</strong> For individuals receiving Supported Living or Family Living, the Agency shall maintain in the individual’s home a complete and current confidential case file for each individual. For individuals receiving Independent Living Services, rather than maintaining this file at the individual’s home, the complete and current confidential case file for each individual shall be maintained at the agency’s administrative site. Each file shall include the following:</td>
<td></td>
</tr>
<tr>
<td>(1) Complete and current ISP and all supplemental plans specific to the individual;</td>
<td>(1) Special Health Care Needs</td>
</tr>
<tr>
<td>(2) Complete and current Health Assessment Tool;</td>
<td>° Meal Time Plan (#3)</td>
</tr>
<tr>
<td>(3) Current emergency contact information, which includes the individual’s address, telephone number, names and telephone numbers of residential Community Living Support providers, relatives, or guardian or conservator, primary care physician's name(s) and telephone number(s), pharmacy name, address and telephone number and dentist name, address and telephone number, and health plan;</td>
<td></td>
</tr>
<tr>
<td>(4) Up-to-date progress notes, signed and dated by the person making the note for at least the past month (older notes may be transferred to the agency office);</td>
<td></td>
</tr>
<tr>
<td>(5) Data collected to document ISP Action Plan implementation</td>
<td></td>
</tr>
<tr>
<td>(6) Progress notes written by direct care staff and by nurses regarding individual health status and physical conditions including action taken in response to identified changes in condition for at least the past month;</td>
<td></td>
</tr>
<tr>
<td>(7) Physician’s or qualified health care providers</td>
<td></td>
</tr>
</tbody>
</table>

Based on record review, the Agency failed to maintain a complete and confidential case file in the residence for 1 of 3 Individuals receiving Supported Living Services. The following was not found, incomplete and/or not current:

- **Special Health Care Needs**
  - Meal Time Plan (#3)
written orders;

(8) Progress notes documenting implementation of a physician’s or qualified health care provider’s order(s);

(9) Medication Administration Record (MAR) for the past three (3) months which includes:
   (a) The name of the individual;
   (b) A transcription of the healthcare practitioners prescription including the brand and generic name of the medication;
   (c) Diagnosis for which the medication is prescribed;
   (d) Dosage, frequency and method/route of delivery;
   (e) Times and dates of delivery;
   (f) Initials of person administering or assisting with medication; and
   (g) An explanation of any medication irregularity, allergic reaction or adverse effect.

(h) For PRN medication an explanation for the use of the PRN must include:
   (i) Observable signs/symptoms or circumstances in which the medication is to be used, and
   (ii) Documentation of the effectiveness/result of the PRN delivered.

(i) A MAR is not required for individuals participating in Independent Living Services who self-administer their own medication. However, when medication administration is provided as part of the Independent Living Service a MAR must be maintained at the individual’s home and an updated copy must be placed in the agency file on a weekly basis.

(10) Record of visits to healthcare practitioners including any treatment provided at the visit and a record of all diagnostic testing for the current ISP year; and

(11) Medical History to include: demographic data, current and past medical diagnoses including the cause (if known) of the developmental disability and any psychiatric diagnosis, allergies (food,
environmental, medications), status of routine adult health care screenings, immunizations, hospital discharge summaries for past twelve (12) months, past medical history including hospitalizations, surgeries, injuries, family history and current physical exam.
<table>
<thead>
<tr>
<th>Tag # 6L25 (CoP)</th>
<th>Residential Health &amp; Safety (Supported Living &amp; Family Living)</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>CHAPTER 6. VIII. COMMUNITY LIVING SERVICE PROVIDER AGENCY REQUIREMENTS</strong></td>
<td></td>
</tr>
<tr>
<td><strong>L. Residence Requirements for Family Living Services and Supported Living Services</strong></td>
<td></td>
</tr>
<tr>
<td>(1) Supported Living Services and Family Living Services providers shall assure that each individual’s residence has:</td>
<td></td>
</tr>
<tr>
<td>(a) Battery operated or electric smoke detectors, heat sensors, or a sprinkler system installed in the residence;</td>
<td></td>
</tr>
<tr>
<td>(b) General-purpose first aid kit;</td>
<td></td>
</tr>
<tr>
<td>(c) When applicable due to an individual’s health status, a blood borne pathogens kit;</td>
<td></td>
</tr>
<tr>
<td>(d) Accessible written procedures for emergency evacuation e.g. fire and weather-related threats;</td>
<td></td>
</tr>
<tr>
<td>(e) Accessible telephone numbers of poison control centers located within the line of sight of the telephone;</td>
<td></td>
</tr>
<tr>
<td>(f) Accessible written documentation of actual evacuation drills occurring at least three (3) times a year. For Supported Living evacuation drills shall occur at least once a year during each shift;</td>
<td></td>
</tr>
<tr>
<td>(g) Accessible written procedures for the safe storage of all medications with dispensing instructions for each individual that are consistent with the Assisting with Medication Administration training or each individual’s ISP; and</td>
<td></td>
</tr>
<tr>
<td>(h) Accessible written procedures for emergency placement and relocation of individuals in the event of an emergency evacuation that makes the residence unsuitable for occupancy. The emergency evacuation procedures shall address, but are not limited to, fire, chemical and/or hazardous waste spills, and flooding.</td>
<td></td>
</tr>
</tbody>
</table>

**Scope and Severity Rating: F**

Based on observation, the Agency failed to ensure that each individual’s residence met all requirements within the standard for 1 of 1 Supported Living residences.

The following items were not found, not functioning or incomplete:

**Supported Living Requirements:**

*Note: Individuals #1, 2 & 3 live in the same residence.*

- Accessible written documentation of actual evacuation drills occurring at least three (3) times a year. For Supported Living evacuation drills shall occur at least once a year during each shift (#1, 2 & 3)

- Accessible written procedures for the safe storage of all medications with dispensing instructions for each individual that are consistent with the Assisting with Medication Administration training or each individual’s ISP (#1, 2 & 3)

- Accessible written procedures for emergency placement and relocation of individuals in the event of an emergency evacuation that makes the residence unsuitable for occupancy. The emergency evacuation procedures shall address, but are not limited to, fire, chemical and/or hazardous waste spills, and flooding (#1, 2 & 3)
ADDITIONAL FINDINGS: Reimbursement Deficiencies

BILLING
TAG #1A12

Developmental Disabilities (DD) Waiver Service Standards effective 4/1/2007 Chapter 1. III. PROVIDER AGENCY DOCUMENTATION OF SERVICE DELIVERY AND LOCATION

B. Billable Units: The documentation of the billable time spent with an individual shall be kept on the written or electronic record that is prepared prior to a request for reimbursement from the HSD. For each unit billed, the record shall contain the following:

1. Date, start and end time of each service encounter or other billable service interval;
2. A description of what occurred during the encounter or service interval; and
3. The signature or authenticated name of staff providing the service.

Billing for Community Living (Supported Living) and Community Inclusion (Adult Habilitation) services was reviewed for 3 of 3 individuals. Progress notes and billing records supported billing activities for the months of March, April, and May, 2010.